

FILE 'HOME' ENTERED AT 14:14:18 ON 16 AUG 2001

```
=> file medline caplus embase biosis
COST IN U.S. DOLLARS          SINCE FILE      TOTAL
                                ENTRY          SESSION
  FULL, ESTIMATED COST        0.15          0.15
```

FILE 'MEDLINE' ENTERED AT 14:14:30 ON 16 AUG 2001

FILE 'CPLUS' ENTERED AT 14:14:30 ON 16 AUG 2001  
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.  
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.  
COPYRIGHT (C) 2001 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'EMBASE' ENTERED AT 14:14:30 ON 16 AUG 2001  
COPYRIGHT (C) 2001 Elsevier Science B.V. All rights reserved.

FILE 'BIOSIS' ENTERED AT 14:14:30 ON 16 AUG 2001  
COPYRIGHT (C) 2001 BIOSIS(R)

=> s MIIC (P) (MHC (3N) Class II)  
L1 155 MIIC (P) (MHC (3N) CLASS II)

```
=> dup rem l1
PROCESSING COMPLETED FOR L1
L2      55 DUP REM L1 (100 DUPLICATES REMOVED)
```

=> s 12 and PD<1999  
'1999' NOT A VALID FIELD CODE  
2 FILES SEARCHED...  
3 FILES SEARCHED...  
L3 14 L2 AND PD<1999

=> dis 13 ibib kwic

L3 ANSWER 1 OF 14 CAPLUS COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 1998:737920 CAPLUS  
 DOCUMENT NUMBER: 130:152130  
 TITLE: The role of the endocytic system in antigen presentation  
 AUTHOR(S): Geuze, Hans J.  
 CORPORATE SOURCE: Laboratory of Cell Biology and Institute of Biomembranes, Utrecht University, Neth.  
 SOURCE: Electron Microsc. 1998, Proc. Int. Congr., 14th (1998), Volume 1, 853-854. Editor(s): Calderon Benavides, Hector A.; Jose Yacaman, Miguel. Institute of Physics Publishing: Bristol, UK.  
 CODEN: 66Y4A  
 DOCUMENT TYPE: Conference; General Review  
 LANGUAGE: English  
 REFERENCE COUNT: 8  
 REFERENCE(S):  
 (1) Glickman, J; J Cell Biol 1996, V132, P769 CAPLUS  
 (3) Kleijmeer, M; J Cell Biol 1997, V139, P639 CAPLUS  
 (4) Kleijmeer, M; Methods Companion Methods Enzymol 1996, V10, P191 CAPLUS  
 (5) Neefjes, J; Cell 1990, V61, P171 CAPLUS  
 (6) Peters, P; Nature 1991, V349, P669 CAPLUS  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

SO Electron Microsc. 1998, Proc. Int. Congr., 14th (1998), Volume 1, 853-854. Editor(s): Calderon Benavides, Hector A.; Jose Yacaman, Miguel. Publisher: Institute of Physics Publishing, Bristol, UK. CODEN: 66YYA4

AB A review and discussion with 8 refs. Endocytosis of antigen varies for different types of antigen-presenting cells (APCs) and can occur by phagocytosis, (macro)pinocytosis or receptor-mediated endocytosis. All entries into the APCs converge at endosomes. The authors discuss the structures collectively called MHC class II compartments) and how these may be involved in antigen processing.

=> dis 13 1-14 ibib kwic

L3 ANSWER 1 OF 14 CAPLUS COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 1998:737920 CAPLUS  
 DOCUMENT NUMBER: 130:152130  
 TITLE: The role of the endocytic system in antigen presentation  
 AUTHOR(S): Geuze, Hans J.  
 CORPORATE SOURCE: Laboratory of Cell Biology and Institute of Biomembranes, Utrecht University, Neth.  
 SOURCE: *Electron Microscopie* 1998, Proc. Int. Congr., 14th (1998), Volume 1, 653-654. Editor(s): Calderon Benavides, Hector A.; Jose Yacaman, Miguel. Institute of Physics Publishing: Bristol, UK.  
 CODEN: 66Y4A  
 DOCUMENT TYPE: Conference; General Review  
 LANGUAGE: English  
 REFERENCE COUNT: 8  
 REFERENCE(S):  
 (1) Glickman, J; *J Cell Biol* 1996, V132, P769 CAPLUS  
 (3) Kleijmeer, M; *J Cell Biol* 1997, V139, P639 CAPLUS  
 (4) Kleijmeer, M; *Methods Companion Methods Enzymol* 1996, V10, P191 CAPLUS  
 (5) Neefjes, J; *Cell* 1990, V61, P171 CAPLUS  
 (6) Peters, P; *Nature* 1991, V349, P669 CAPLUS

ALL CITATIONS AVAILABLE IN THE RE FORMAT  
SO Electron Microsc. 1998, Proc. Int. Congr., 14th (1998), Volume  
1, 853-854. Editor(s): Calderon Benavides, Hector A.; Jose Yacaman,  
Miguel. Publisher: Institute of Physics Publishing, Bristol, UK.  
S0000-0000/98/0001-0853\$15.00  
© 1998 IOP Publishing Ltd

AB CODEN: 66YYA4  
A review and discussion with 8 refs. Endocytosis of antigen varies for different types of antigen-presenting cells (APCs) and can occur by phagocytosis, (macro)pinocytosis or receptor-mediated endocytosis. All entries into the APCs converge at endosomes. The authors discuss the structures collectively called MHC class II compartments) and how these may be involved in antigen processing.

L3 ANSWER 2 OF 14 CAPLUS COPYRIGHT 2001 ACS  
ACCESSION NUMBER: 1998:243907 CAPLUS

DOCUMENT NUMBER: 129:26741  
TITLE: Multiple signals regulate the intracellular trafficking of HLA-DM in B-lymphoblastoid cells  
AUTHOR(S): Copier, J.; Potter, P.; Sacks, S. H.; Kelly, A. P.  
CORPORATE SOURCE: Department of Nephrology and Transplantation, Guy's Hospital, London, UK  
SOURCE: Immunology (1998), 93(4), 505-510  
PUBLISHER: Blackwell Science Ltd.  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
SO Immunology (1998), 93(4), 505-510  
CODEN: IMMUAM; ISSN: 0019-2805  
IT Organelle (MIIC (MHC class II compartment); regulation of intracellular trafficking of HLA-DM in B-cells)

L3 ANSWER 3 OF 14 CAPLUS COPYRIGHT 2001 ACS  
ACCESSION NUMBER: 1997:1443696 CAPLUS  
DOCUMENT NUMBER: 127:175103  
TITLE: Decreased endosomal delivery of major histocompatibility complex class II-invariant chain complexes in dynamin-deficient cells  
AUTHOR(S): Wang, Kena; Peterson, Per A.; Karlsson, Lars R. W. Johnson Pharmaceutical Research Institute, San Diego, CA, 92121, USA  
CORPORATE SOURCE: J. Biol. Chem. (1997), 272(27), 17055-17060  
SOURCE: CODEN: JBCHA3; ISSN: 0021-9258  
PUBLISHER: American Society for Biochemistry and Molecular Biology  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
SO J. Biol. Chem. (1997), 272(27), 17055-17060  
CODEN: JBCHA3; ISSN: 0021-9258  
IT Organelle (MIIC (MHC class II compartment); invariant chain/MHC class II sorting is dynamin-dependent)

L3 ANSWER 4 OF 14 CAPLUS COPYRIGHT 2001 ACS  
ACCESSION NUMBER: 1997:265045 CAPLUS  
DOCUMENT NUMBER: 127:16339  
TITLE: Assembly of an abundant endogenous major histocompatibility complex class II/peptide complex in class II compartments  
AUTHOR(S): Morkowski, Stanislaw; Raposo, Graca; Kleijmeer, Monique; Geuze, Hans J.; Rudensky, Alexander Y.  
CORPORATE SOURCE: School Medicine, University Washington, Seattle, WA, 98195, USA  
SOURCE: Eur. J. Immunol. (1997), 27(3), 609-617  
CODEN: EJIMAF; ISSN: 0014-2980  
PUBLISHER: VCH  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
SO Eur. J. Immunol. (1997), 27(3), 609-617  
CODEN: EJIMAF; ISSN: 0014-2980  
ST peptide MHC class II assembly lymphocyte; B cell MIIC compartment peptide MHC  
IT Organelle (MIIC (MHC class II compartment); endogenous major histocompatibility complex class II/peptide complex assembled in)

L3 ANSWER 5 OF 14 CAPLUS COPYRIGHT 2001 ACS  
ACCESSION NUMBER: 1996:548779 CAPLUS  
DOCUMENT NUMBER: 125:216300  
TITLE: Characterization of MHC class II compartments by immunoelectron microscopy  
AUTHOR(S): Kleijmeer, Monique J.; Raposo, Graca; Geuze, Hans J.  
CORPORATE SOURCE: Dep. Cell Biology, Utrecht Univ., Utrecht, 3584 CX, Neth.  
SOURCE: Methods (San Diego) (1996), 10(2), 191-207  
CODEN: MTHDE9; ISSN: 1046-2023  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
SO Methods (San Diego) (1996), 10(2), 191-207  
CODEN: MTHDE9; ISSN: 1046-2023

AB At present the best way to det. the precise intracellular localization of proteins, in a potentially semiquant. way, is the combination of ultrathin cryosectioning and immunogold labeling. This paper focuses on the intracellular localization of MHC class II mols., which are involved in the T helper response to exogenous antigens. Newly synthesized MHC class II heterodimers assoc. with invariant chain mols., which in turn direct the MHC class II complex to the endocytic route. Proteolytic digestion of the invariant chain frees MHC class II mols. so that they can bind antigenic peptides. Immunoelectron microscopy has been an important tool to identify the endocytic compartments that are enriched in MHC class II and that are the potential sites of antigenic peptide binding. The methods that can be used to characterize MHC class II compartments (MIICs) in various antigen-presenting cells (APCs) are described in detail. In all APCs studies so far, MIICs are situated late in the endocytic pathway and display lysosomal characteristics. Still, immunoelectron microscopy allows us to define subsets of MIICs, which can be distinguished by their morphol., accessibility to endocytic tracers, and expression of invariant chain and HLA-DM. Different types of MIICs can be found that display internal vesicles (multivesicular), internal membrane sheets (multilaminar), or both. The multivesicular type of MIIC contains detectable invariant chain and is the primary site of antigen entry. The multilaminar MIIC is situated later in the endocytic route and has lost most of the invariant chain antigenicity. These data suggest a sequential maturation of MIICs, which correlates with the degrdn. of invariant chain and the subsequent binding of antigenic peptides.

L3 ANSWER 6 OF 14 CAPLUS COPYRIGHT 2001 ACS  
ACCESSION NUMBER: 1995:858363 CAPLUS  
DOCUMENT NUMBER: 123:253975  
TITLE: A lysosomal targeting signal in the cytoplasmic tail

of the  $\beta$  chain di HLA-DM to MHC class II compartments  
AUTHOR(S): Marks, Michael S.; Roche, Paul A.; van Donselaar, compartments  
Elly; Woodruff, Lauren; Peters, Peter J.; Bonifacino, Juan S.  
CORPORATE SOURCE: Cell Biology and Metabolism Branch, Natl. Inst.  
Health, Bethesda, MD, 20892, USA  
SOURCE: J. Cell Biol. (1995), 131(2), 351-69  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
SO J. Cell Biol. (1995), 131(2), 351-69  
CODEN: JCLBA3; ISSN: 0021-9525  
IT Organelle (MIIC (MHC class II compartment); lysosomal targeting signal in cytoplasmic tail of  $\beta$  chain directs HLA-DM to MHC class II compartments)

L3 ANSWER 7 OF 14 CAPLUS COPYRIGHT 2001 ACS  
ACCESSION NUMBER: 1995:328971 CAPLUS  
DOCUMENT NUMBER: 122:103518  
TITLE: How MHC class II molecules reach the endocytic pathway  
AUTHOR(S): Benaroch, Philippe; Yilla, Mamadi; Raposo, Graca; Ito, Kouichi; Miwa, Kiyoshi; Geuze, Hans J.; Ploegh, Hidde L.  
CORPORATE SOURCE: Center Cancer Research, Dep. Biology, Massachusetts Institute Technology, Cambridge, MA, 02139, USA  
SOURCE: EMBO J. (1995), 14(1), 37-49  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
SO EMBO J. (1995), 14(1), 37-49  
CODEN: EMJODG; ISSN: 0261-4189  
IT Organelle (MIIC (MHC class II compartment); endocytic trafficking of MHC class II/invariant chain complexes in human B-cells)

L3 ANSWER 8 OF 14 CAPLUS COPYRIGHT 2001 ACS  
ACCESSION NUMBER: 1994:532185 CAPLUS  
DOCUMENT NUMBER: 121:132185  
TITLE: A novel lysosomal compartment engaged in antigen presentation  
AUTHOR(S): Geuze, Hans  
CORPORATE SOURCE: Laboratory Cell Biology, Utrecht University School Medicine, Utrecht, Neth.  
SOURCE: Eur. J. Cell Biol. (1994), 64(1), 3-6  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
SO Eur. J. Cell Biol. (1994), 64(1), 3-6  
CODEN: EJCBDN; ISSN: 0171-9335  
AB The authors attempted to identify the endocytic compartments involved in antigen processing and peptide binding to MHC class II. Using immunogold labeling of ultrathin cryosections, MHC class II, invariant chain, and organelle markers were localized in a variety of antigen-presenting cells. In human B-cell the majority of MHC class II mols. was found in a compartment called the MHC class II-enriched compartment (MIIC) with a characteristic morphol.: it contains internal vesicles and membrane sheets. MIIC in B-cells were shown to share several features with lysosomes.

L3 ANSWER 9 OF 14 CAPLUS COPYRIGHT 2001 ACS  
ACCESSION NUMBER: 1994:506199 CAPLUS  
DOCUMENT NUMBER: 121:106199  
TITLE: Major histocompatibility complex class II molecules induce the formation of endocytic MIIC-like structures  
AUTHOR(S): Calafat, Jero; Nijenhuis, Marga; Janssen, Hans; Tulp, Abraham; Dusseljee, Simone; Wubbolts, Richard; Neefjes, Jacques  
CORPORATE SOURCE: Division Cellular Biochem., Netherlands Cancer Inst., Amsterdam, 1066 CX, Neth.  
SOURCE: J. Cell Biol. (1994), 126(4), 967-77  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
SO J. Cell Biol. (1994), 126(4), 967-77  
CODEN: JCLBA3; ISSN: 0021-9525  
AB During biosynthesis, major histocompatibility complex class II mols. are transported to the cell surface through a late endocytic multilaminar structure with lysosomal characteristics. This structure did not resemble any of the previously described endosomal compartments and was termed MIIC (for MHC class II compartment). The authors show here that continuous protein synthesis is required for the maintenance of MIIC in B cells. Transfection of class II mols. in human embryonal kidney cells induces the formation of multilaminar endocytic structures that are morphol. analogous to MIIC in B cells. Two lysosomal proteins (CD63 and lamp-1), which are expressed in MIIC of B cells, are also present in the structures induced by expression of major histocompatibility complex class II mols. Moreover, endocytosed HRP enters the induced structures defining them as endocytic compartments. Exchanging the transmembrane and cytoplasmic tail of the class II  $\alpha$  and  $\beta$  chains for that of HLA-B27 does not result in the induction of multilaminar structures, and the chimeric class II mols. are now located in multivesicular structures. Thus, expression of class II mols. is sufficient to induce the formation of characteristic MIIC-like multilaminar structures.

L3 ANSWER 10 OF 14 CAPLUS COPYRIGHT 2001 ACS  
ACCESSION NUMBER: 1994:320969 CAPLUS  
DOCUMENT NUMBER: 120:320969  
TITLE: Antigen processing and class II MHC peptide-loading compartments in human B-lymphoblastoid cells  
AUTHOR(S): West, Michele A.; Lucocc, John M.; Watts, Colin  
CORPORATE SOURCE: Med. Sci. Inst., Univ. Dundee, Dundee, DD1 4HN, UK  
SOURCE: Nature (London) (1994), 369(6476), 147-51  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
SO Nature (London) (1994), 369(6476), 147-51

CODEN: NATUAS; ISSN: 0028-0836  
 IT Organelle  
 (MHC class II-assocd.  
 compartment), of human B-cells, in antigen processing and MHC  
 class II complex loading with peptide)

L3 ANSWER 11 OF 14 CAPLUS COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 1993:536899 CAPLUS  
 DOCUMENT NUMBER: 119:136699  
 TITLE: The molecular basis for T cell help in humoral immunity: CD40 and its ligand, gp39  
 AUTHOR(S): Marshall, Lisa S.; Aruffo, Alejandro; Ledbetter, Jeffrey A.; Noelle, Randolph J.  
 CORPORATE SOURCE: Dartmouth Med. Sch., Lebanon, NH, 03756, USA  
 SOURCE: J. Clin. Immunol. (1993), 13(3), 165-74  
 CODEN: JCIMD; ISSN: 0271-9142  
 DOCUMENT TYPE: Journal: General Review  
 LANGUAGE: English  
 SO J. Clin. Immunol. (1993), 13(3), 165-74  
 CODEN: JCIMD; ISSN: 0271-9142  
 AB A review and discussion with 73 refs. Thymus-dependent (TD) humoral immune responses require cognate interactions between B cells and CD4+ helper T cells (Th). Since TD antigens do not express highly repeated epitopes, the binding of antigen to membrane IgM and membrane IgD (mIg) is insufficient to trigger B cell cycle entry and subsequent antibody prodn. Although incapable of directly triggering B cell activation, once bound to mIg, TD antigen is endocytosed and processed by antigen-specific B cells. The processed antigen is expressed on the B cell surface in a complex with MHC class II mol., and presented for Th recognition. Ligation of CD4 and the T cell receptor (TcR) by the antigen/MHC class II complex, together with the interaction of other surface mol. ligand-receptor pairs, including CD28-B7, LFA1-ICAM1, and CD4-MHC class II, activates the Th. Once activated, Th rapidly express lymphokine genes and a membrane protein, gp39, which is essential for the reciprocal activation of the cognate, antigen-presenting B cell. The interaction of gp39 with its receptor CD40, on the B cell, derives B cell cycle entry and induces B cell responsiveness to the growth and differentiative effects of lymphokines.

L3 ANSWER 12 OF 14 BIOSIS COPYRIGHT 2001 BIOSIS  
 ACCESSION NUMBER: 1998:363067 BIOSIS  
 DOCUMENT NUMBER: PREV199800363067  
 TITLE: Donor pretreatment with Flt-3 ligand augments antidor cytotoxic T lymphocyte, natural killer, and lymphokine-activated killer cell activities within liver allografts and alters the pattern of intragraft apoptotic activity.  
 AUTHOR(S): Qian, Shiguang (1); Lu, Lina; Fu, Fumin; Li, Wei; Pan, Fan; Steptoe, Raymond J.; Chambers, Frances G.; Starzl, Thomas E.; Fung, John J.; Thomson, Angus W. (1)  
 CORPORATE SOURCE: (1) W1540 Biomed. Sci. Tower, Univ. Pittsb. Med. Cent., 200 Lothrop St., Pittsburgh, PA 15213 USA  
 SOURCE: Transplantation (Baltimore), (June 27, 1998) Vol. 65, No. 12, pp. 1590-1598.  
 ISSN: 0041-1337.  
 DOCUMENT TYPE: Article  
 LANGUAGE: English  
 SO Transplantation (Baltimore), (June 27, 1998) Vol. 65, No. 12, pp. 1590-1598.  
 ISSN: 0041-1337.  
 AB . . . cytokine that strikingly augments functional dendritic cells (DCs) within lymphoid and nonlymphoid tissue. Methods. The expression of costimulatory molecules and MHC class II antigen on DCs isolated from livers of FL-treated B10 (H2b) mice (10 mug/day; 10 days) was examined by flow cytometric. . . in primary mixed leukocyte cultures. B10 livers from FL-treated donors were transplanted orthotopically into naive C3H (H2k) recipients. Donor cells (MHC class (II+)) in recipient spleens were identified by immunohistochemistry. Antidor cytotoxic T lymphocyte activity, and both natural killer and lymphokine-activated . . .

L3 ANSWER 13 OF 14 BIOSIS COPYRIGHT 2001 BIOSIS  
 ACCESSION NUMBER: 1997:96196 BIOSIS  
 DOCUMENT NUMBER: PREV199799395399  
 TITLE: Egress of newly peptide-loaded MHC class II molecules from the MHC to the plasma membrane is independent of early endosomes.  
 AUTHOR(S): Pond, Leslie; Watts, Colin  
 CORPORATE SOURCE: Dep. Biochemistry, Med. Sciences Inst., Univ. Dundee, Dundee DD1 4HN UK  
 SOURCE: Molecular Biology of the Cell, (1996) Vol. 7, No. SUPPL., pp. 325A.  
 Meeting Info.: Annual Meeting of the 6th International Congress on Cell Biology and the 36th American Society for Cell Biology San Francisco, California, USA December 7-11, 1996  
 ISSN: 1059-1524.  
 DOCUMENT TYPE: Conference; Abstract; Conference  
 LANGUAGE: English  
 TI Egress of newly peptide-loaded MHC class II molecules from the MHC to the plasma membrane is independent of early endosomes.  
 SO Molecular Biology of the Cell, (1996) Vol. 7, No. SUPPL., pp. 325A.  
 Meeting Info.: Annual Meeting of the 6th International Congress on Cell Biology and the 36th American Society for Cell Biology San Francisco, California, USA December 7-11, 1996  
 ISSN: 1059-1524.

L3 ANSWER 14 OF 14 BIOSIS COPYRIGHT 2001 BIOSIS  
 ACCESSION NUMBER: 1997:53591 BIOSIS  
 DOCUMENT NUMBER: PREV199799352794  
 TITLE: In vitro differentiation of CD34+ hematopoietic progenitor cells towards distinct dendritic cell subsets of the MHC-positive Langerhans cell- and the interdigitating dendritic cell type.  
 AUTHOR(S): Lindemann, Albrecht (1); Koehler, Gabriele; Mackensen, Andreas (1); Veelken, Hendrik (1); Rosenthal, Felicia M. (1); Schaefer, Hans Eckhart; Fisch, Paul (1); Mertelsmann, Roland (1); Herbst, Birgit (1)  
 CORPORATE SOURCE: (1) Dep. Med. I, Univ. Med. Cent., Freiburg Germany  
 SOURCE: Blood, (1996) Vol. 88, No. 10 SUPPL. 1 PART 1-2, pp. 153A.  
 Meeting Info.: Thirty-eighth Annual Meeting of the American

Society of Hematology Orlando, Florida, USA December 6-10,  
1996

ISSN: 0006-4971.

DOCUMENT TYPE: Conference; Abstract; Conference

LANGUAGE: English

SO Blood, (1996) Vol. 88, No. 10 SUPPL. 1 PART 1-2, pp. 153A.  
Meeting Info.: Thirty-eighth Annual Meeting of the American Society of  
Hematology Orlando, Florida, USA December 6-10, 1996

ISSN: 0006-4971.

IT GRANULOCYTES; IN-VITRO; MACROPHAGES; MAJOR HISTOCOMPATIBILITY  
COMPLEX-CLASS II COMPARTMENT-POSITIVE INTERDIGITATING DENDRITIC CELL  
TYPE; MAJOR HISTOCOMPATIBILITY COMPLEX-CLASS II COMPARTMENT-POSITIVE  
LANGERHANS CELL TYPE; MHC-CLASS II  
COMPARTMENT-POSITIVE INTERDIGITATING DENDRITIC CELL TYPE; MHC  
-CLASS II COMPARTMENT-POSITIVE LANGERHANS CELL  
TYPE; MHC-POSITIVE INTERDIGITATING DENDRITIC CELL TYPE;  
MHC-POSITIVE LANGERHANS CELL TYPE; MONOCYTES; T CELL  
ACTIVATION; T CELL TRAFFICKING

=> end

ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF

LOGOFF? (Y)/N/HOLD:y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
	38.84	38.99

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-3.53	-3.53

STN INTERNATIONAL LOGOFF AT 14:20:21 ON 16 AUG 2001